Hill AFB Basewide QAPP

Revision: 0.0 Date: July 1999

8.0 INTERNAL QUALITY CONTROL CHECKS

8.0.0.1. Internal quality control checks are used to evaluate whether method performance

is within acceptable limits of precision and accuracy. The following sections describe the

internal QC that shall be followed for both field and laboratory activities.

8.1 FIELD SAMPLE COLLECTION

8.1.0.1. Quality control samples are used to assess field sampling techniques and

environmental conditions during sample collection and transportation and typically

include (but are not limited to) field duplicates, trip blanks, ambient air blanks, source

water blanks, and equipment blanks. Quality control sample requirements for field

samples for both screening and definitive data are described in Table 3-6.

8.2 FIELD MEASUREMENTS

8.2.0.1. QC procedures for field measurements shall include checking the reproducibility

of the measurements by obtaining multiple readings on a single grab sample (if practical)

or back checking meter accuracy against a standard, calibrating the field instruments or

equipment as discussed in SOP 2 and monitoring instrument calibration throughout each

day of sampling.

8.3 LABORATORY ANALYSIS

8.3.0.1. The general objectives of the laboratory QC program are to:

• Ensure that all procedures are documented, including any changes in

administrative and/or technical procedures.

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• Ensure that all analytical procedures are validated and conducted according to

method guidelines and laboratory SOPs.

• Monitor the performance of the laboratory using a systematic inspection

program.

• Ensure that all data are properly reported and archived.

8.3.0.2. Contract laboratories shall conduct internal quality control checks for analytical

methods in accordance with their SOPs, the individual method requirements, and this

QAPP. The laboratory shall notify the Prime Contractor's Project Manager in writing

before making significant changes to the project-specific work plans, this QAPP or

analytical methodology.

8.3.0.3. Laboratory quality control consists of two distinct components, a laboratory

component and a matrix component. The laboratory component measures the

performance of the laboratory analytical process during sample analyses, while the matrix

component measures the effects of a specific media on the method performance.

The QC samples that shall be used to assess the laboratory component and the media

component of analysis are listed in Tables 3-4 and 3-6. The criteria against which the QC

data shall be evaluated are listed in the quality control criteria tables included in

Appendices A through H. Corrective actions for instrument calibrations or out of

compliance QC sample data are listed in the corrective action summary tables, and also

are included in Appendices A through H.

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